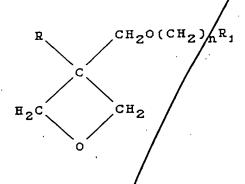
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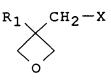
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1. A method of making a mono-substituted fluorinated oxetane (FOX) monomer having the structure:



where n is 1 to 3, R is methyl or ethyl, and R_f is linear or branched chain fluorinated alkyl and isoalkyl having from 1 to 20 carbons or oxa-perfluorinated polyether, having from 4 to about 60 carbons comprising the steps of:

a) providing a mono-substituted oxetane premonomer having the structure:



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where R_1 is selected from the group consisting of methyl and ethyl and X is a leaving group selected from the group consisting of bromo, chloro, iodo and aryl sulfonate, said premonomer being dissolved in a solvent to provide a premonomer solution;

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b) charging a reaction vessel with an aqueous solution of said mono-substituted oxetane premonomer, a fluoroalcohol, a

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phase transfer catalyst and a strong base; and

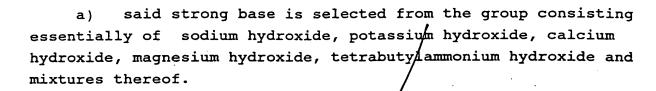
- c) heating said solution to a temperature of 80-85°C until reaction is complete to form the FOX monomer as a separate organic layer.
- 2. A method of making a mono-substituted FOX monomer as in claim 1 which includes the steps of:
 - a) cooling the reaction mixture; and
- b) separating the mono-substituted FOX monomer as an organic layer from the aqueous reaction mixture.
- 3. A method of making a mono-substituted fluorinated oxetane monomer as in claim 1 wherein.
- a) said fluorinated alcohol is selected from the group consisting essentially of trifluoroethanol, heptafluorobutanol, pentadecafluorooctanol, tridecafluorooctanol, other fluorinated alcohols having the following formulas:
 - a) $HO(CH_2)/(CF_2)_X-F$;
 - b) $HQCH_2CF_2(OCF_2CF_2)_X-F$

wherein n is 1 to β and x is 1 to 20 and mixtures thereof.

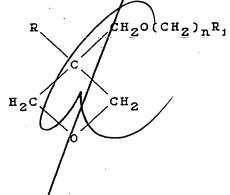
- 4. A method of making a mono-substituted FOX monomer as in claim 3 wherein:
- a) said phase transfer catalyst is selected from the group consisting essentially of tetrabutylammonium bromide, tetraethylammonium bromide, tetratmethylammonium iodide, cetyltributylammonium bromide, crown ethers, glycols and mixtures thereof.
- 5. A method of making a mono-substituted FOX monomer as in claim 4 wherein:

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- 6. A method of making a mono-substituted FOX as in claim 5 wherein:
- a) said strong base is potassium hydroxide and said phase transfer catalyst is tetrabutylammonium bromide, and said temperature is in the range of from about 80°C to about 85°C.
 - 7. A mono-substituted fluorinated oxetane monomer having the structure:



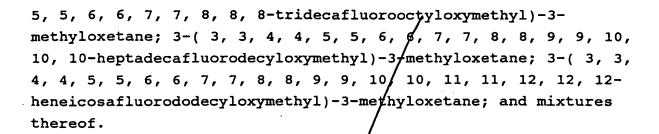
Where:

n is 1/to 3;

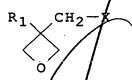
R is methyl or ethyl; and R_f is linear or branched chain fluorinated alkyl and isoalkyl having from 1 to 20 carbons or oxaperfluorinated polyether, having from 4 to about 60 carbons.

8. A mono-substituted fluorinated oxetane monomer as in claim 7 including 3-(2,2,2-trifluoroethoxymethyl)-3-methyloxetane; 3-(2,2,3,3,4,4,4-heptafluorobutoxymethyl)-3-methyloxetane; 3-(2,2,3,3,4,4,5,5,6,6,7,7,8,8,8-pentadecafluorooctyloxymethyl)-3-methyloxetane; 3-(3,3,4,4,

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- 9. A mono-substituted fluorinated oxetane (FOX) monomer produced by the process comprising the steps of:
- a) providing a mono-substituted oxetane premonomer having the structure:

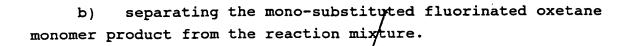


where R₁ is selected from the group consisting of methyl and ethyl and X is a leaving group selected from the group consisting of bromo, chloro, iodo and aryl sulfonate, said premonomer being diluted in a solvent to provide a premonomer solution;

- b) suspending a dispersion of a strong base in an aprotic solvent to provide a strong base suspension;
- c) adding a fluor nated alcohol to said strong base suspension to produce a fluorinated alkoxide solution; and
- d) adding said premonomer solution to said fluorinated alkoxide while heating the reaction mixture to a temperature of about 50 to about 125°C to permit a displacement reaction whereby said fluorinated alkoxide displaces said leaving group to produce the mono-substituted fluorinated oxetane monomer.
- 10. A mono-substituted FOX monomer produced by the process as in claim 9 which includes the steps of:
- a) quenching the displacement reaction upon consumption of the starting materials; and

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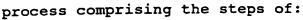
- 11. A mono-substituted FOX monomer produced by the process as in claim 9 wherein:
- a) said fluorinated alcohol is selected from the group consisting essentially of trifluoroethanol, heptafluorobutanol, pentadecafluorooctanol, tridecafluorooctanol, other fluorinated alcohols having the following formulas:
 - a) $HO(CH_2)_n(CF_2)\sqrt{-F}$
 - b) HOCH₂CF₂(OCF₂CF₂)_x-F
 - C) HOCH₂CF(OCF₂CF)_x-F; F₃C CF₃

wherein h is 1 to 3 and x is 1 to 20 and mixtures thereof.

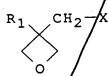
- 12. A mono-substituted FOK monomer produced by the process as in claim 11 wherein:
- a) said strong base is selected from the group consisting essentially of sodium hydride, potassium hydride, potassium t-butoxide, calcium hydride, sodium hydroxide, potassium hydroxide, NaNH₂, n-butyl lithium and lithium diisopropylamide.
- 13. A mono-substituted FOX monomer produced by the process as in claim 12 wherein:
- a) said solvent is selected from the group consisting essentially of dimethylformamide (DMF), dimethylacetamide, DMSO, hexamethylene phosphoramide (HMPA) and mixtures thereof.
- 14. A mono-substituted FOX monomer produced by the process as in claim 13 wherein:
 - a) said temperature is from about 75 to about 85°C.
- 15. A mono-substituted fluorinated monomer produced by the

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a) providing a mono-substituted oxetane premonomer having the structure:



where R_1 is selected from the group consisting of methyl and ethyl and X is a leaving group selected from the group consisting of bromo, chloro, iodo and aryl sulfonate, said premonomer being dissolved in a solvent to provide a premonomer solution;

- b) charging a reaction vessel with an aqueous solution of said mono-substituted oxetane premonomer, a fluoroalcohol, a phase transfer catalyst and a strong base;
- c) heating said solution to a temperature of 80-85°C until reaction is complete to form the FOX monomer as a separate organic layer;
 - d) cooling the reaction mixture; and
- e) separating the mono-substituted fluorinated oxetane monomer as an organic layer from the aqueous reaction mixture.
- 16. A mono-substituted fluorinated monomer produced by the process of claim 15 wherein:
- a) said phase transfer catalyst is selected from the group consisting essentially of tetrabutylammonium bromide, tetraethylammonium bromide, trimethylbutylammonium bromide, tetratmethylammonium iodide, cetyltributylammonium bromide, crown ethers, glycols and mixtures thereof.
- 17. A mono-substituted fluorinated monomer produced by the process of claim 16 wherein:
- a) said fluorinated alcohol is selected from the group consisting essentially of trifluoroethanol, heptafluorobutanol, pentadecafluoroctanol, tridecafluoroctanol, other fluorinated

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alcohols having the following formulas:

- a) $HO(CH_2)_n(CF_2)_x-F$
- b) HOCH₂CF₂(OCF₂CF₂)_x/F
- c) HOCH2CF(OCF2CF)/-F

wherein n is 1 to about 3 and x is 1 to about 20 and mixtures thereof.

- 18. A mono-substituted fluorinated monomer produced by the process of claim 17 wherein:
- a) said strong base is selected from the group consisting essentially of sodium hydroxide and potassium hydroxide, calcium hydroxide, magnesium hydroxide, tetrabutylammonium hydroxide and mixtures thereof.
- 19. A mono-substituted fluorinated monomer produced by the process of claim 18 wherein:
- a) said strong base is potassium hydroxide and said phase transfer catalyst is tetrabutylammonium bromide.